

### AMENDMENTS TO THE CLAIMS

After entering the Substitute Specification, please amend claims 1, 5, and 8, and add new claims 17-20, as provided in the following listing of claims, which will replace all prior versions and listings of claims in the application. Please cancel claims 6, 7, and 13-15 without prejudice to their pursuit in an appropriate continuation or divisional application.

1 (currently amended).      A method ~~to produce~~ of producing one or more cDNA molecules comprising:

- (a)      contacting a sample comprising a cell or a virus with a solid medium, wherein:
  - (i)      the cell or the virus comprises mRNA and genomic DNA;
  - (ii)     the mRNA comprises an mRNA -template  
~~with a solid medium, of interest; and~~
  - (iii)    wherein the solid medium comprises:
    - (i)- a matrix; and
    - (ii)- a composition for inhibiting degradation of the mRNA  
template, wherein:
      - the composition is sorbed to the matrix; and
      - the composition comprises a detergent or surfactant;
- (b)      sorbing at least a portion of the mRNA template to the solid medium;
- (c)      eluting the mRNA from the solid medium while retaining the genomic DNA;  
and
- (ed)     contacting the ~~template mRNA~~ with one or more reverse  
transcriptases under conditions sufficient to synthesize one or more cDNA  
molecules complementary to all or a portion of the mRNA template of interest.

2 (previously presented).      The method of claim 1, wherein the cDNA is a cDNA library.

3 (previously presented).      The method of claim 1, wherein the mRNA is removed from the solid medium prior to the cDNA synthesis.

4 (previously presented). The method of claim 1, wherein the cDNA is double-stranded.

5 (currently amended). The method of claim 1, further comprising:

(de) amplifying the cDNA.

6. - 7. (canceled)

8 (currently amended). The method of claim 1, wherein ~~the matrix contains a composition for substantially inhibiting degradation of the mRNA template, the detergent or surfactant of the composition is an anionic detergent or surfactant and wherein the composition comprising~~further comprises:

(a) a base; and

(b) a chelating agent; ~~and~~

(c) ~~an anionic detergent or surfactant.~~

9 (previously presented). The method of claim 8, wherein the composition further comprises uric acid or a urate salt.

10 (previously presented). The method of claim 1, wherein the matrix comprises a cellulose-based matrix or paper, or a micromesh of synthetic plastic material.

11 (previously presented). The method of claim 1, wherein the matrix is selected from the group consisting of nitrocellulose, cellulose, diazocellulose, carboxymethylcellulose, hydrophilic polymers, polytetra-fluoro-ethylene, fiberglass, porous ceramics, polystyrene, polyvinylchloride, polypropylene, polyethylene, dextran, agarose, agar, starch, and nylon.

12 (previously presented). The method of claim 1, wherein the sample comprising the mRNA template is selected from the group consisting of cells, viruses, viral plaques, and preparations from biological materials.

13 - 15 (canceled).

16 (previously presented). The method of claim 1, wherein the sample comprising the mRNA template is selected from the group consisting of cells, viruses, viral plaques, and preparations from biological materials.

17 (new). The method of claim 8, wherein:

- a. the anionic detergent or surfactant comprises sodium dodecyl sulfate (SDS);
- b. the base comprises Tris or tris-hydroxymethyl methane; and
- c. the chelating agent comprises ethylene diamine tetra-acetic acid (EDTA).

18 (new). The method of claim 1, wherein the cell is a eukaryotic cell.

19 (new). The method of claim 5, wherein the amplifying step comprises contacting at least one cDNA strand with a polymerase under conditions to synthesize one or more cDNA molecules complementary to all or a portion of the template.

20 (new). The method of claim 5, wherein the amplifying step comprises a polymerase chain reaction (PCR).